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Aegilops tauschii draft genome sequence reveals a gene repertoire for wheat adaptation

Jia, Jizeng ; Zhao, Shancen ; et al

Abstract: About 8,000 years ago in the Fertile Crescent, a spontaneous hybridization of the wild diploid grass *Aegilops tauschii* ($2n = 14$; DD) with the cultivated tetraploid wheat *Triticum turgidum* ($2n = 4x = 28$; AABB) resulted in hexaploid wheat (*T. aestivum*; $2n = 6x = 42$; AABBDD). Wheat has since become a primary staple crop worldwide as a result of its enhanced adaptability to a wide range of climates and improved grain quality for the production of baker's flour. Here we describe sequencing the *Ae. tauschii* genome and obtaining a roughly 90-fold depth of short reads from libraries with various insert sizes, to gain a better understanding of this genetically complex plant. The assembled scaffolds represented 83.4% of the genome, of which 65.9% comprised transposable elements. We generated comprehensive RNA-Seq data and used it to identify 43,150 protein-coding genes, of which 30,697 (71.1%) were uniquely anchored to chromosomes with an integrated high-density genetic map. Whole-genome analysis revealed gene family expansion in *Ae. tauschii* of agronomically relevant gene families that were associated with disease resistance, abiotic stress tolerance and grain quality. This draft genome sequence provides insight into the environmental adaptation of bread wheat and can aid in defining the large and complicated genomes of wheat species.

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Aegilops tauschii draft genome sequence reveals a gene repertoire for wheat adaptation

Jizeng Jia^{1*}, Shancen Zhao^{2,3*}, Xiuying Kong^{1*}, Yingrui Li^{2*}, Guangyao Zhao^{1*}, Weiming He^{2*}, Rudi Appels^{4*}, Matthias Pfeifer⁵, Yong Tao², Xueyong Zhang¹, Ruilian Jing¹, Chi Zhang², Youzhi Ma¹, Lifeng Gao¹, Chuan Gao², Manuel Spannagl⁵, Klaus F. X. Mayer⁵, Dong Li², Shengkai Pan², Fengya Zheng^{2,3}, Qun Hu⁶, Xianchun Xia¹, Jianwen Li², Qinsi Liang², Jie Chen², Thomas Wicker⁷, Caiyun Gou², Hanhui Kuang⁶, Genyun He², Yadan Luo², Beat Keller⁷, Qiuju Xia², Peng Lu², Junyi Wang², Hongfeng Zou², Rongzhi Zhang¹, Junyang Xu², Jinlong Gao², Christopher Middleton⁷, Zhiwu Quan², Guangming Liu⁸, Jian Wang², International Wheat Genome Sequencing Consortium†, Huanming Yang², Xu Liu¹, Zhonghu He^{1,9}, Long Mao¹ & Jun Wang^{2,10,11}

About 8,000 years ago in the Fertile Crescent, a spontaneous hybridization of the wild diploid grass *Aegilops tauschii* ($2n = 14$; DD) with the cultivated tetraploid wheat *Triticum turgidum* ($2n = 4x = 28$; AABB) resulted in hexaploid wheat (*T. aestivum*; $2n = 6x = 42$; AABBDD)^{1,2}. Wheat has since become a primary staple crop worldwide as a result of its enhanced adaptability to a wide range of climates and improved grain quality for the production of baker's flour². Here we describe sequencing the *Ae. tauschii* genome and obtaining a roughly 90-fold depth of short reads from libraries with various insert sizes, to gain a better understanding of this genetically complex plant. The assembled scaffolds represented 83.4% of the genome, of which 65.9% comprised transposable elements. We generated comprehensive RNA-Seq data and used it to identify 43,150 protein-coding genes, of which 30,697 (71.1%) were uniquely anchored to chromosomes with an integrated high-density genetic map. Whole-genome analysis revealed gene family expansion in *Ae. tauschii* of agronomically relevant gene families that were associated with disease resistance, abiotic stress tolerance and grain quality. This draft genome sequence provides insight into the environmental adaptation of bread wheat and can aid in defining the large and complicated genomes of wheat species.

We selected *Ae. tauschii* accession AL8/78 for genome sequencing because it has been extensively characterized genetically (Supplementary Information). Using a whole genome shotgun strategy, we generated 398 Gb of high-quality reads from 45 libraries with insert sizes ranging from 200 bp to 20 kb (Supplementary Information). A hierarchical, iterative assembly of short reads employing the parallelized sequence assembler SOAPdenovo³ achieved contigs with an N50 length (minimum length of contigs representing 50% of the assembly) of 4,512 bp (Table 1). Paired-end information combined with an additional 18.4 Gb of Roche/454 long-read sequences was used sequentially to generate 4.23-Gb scaffolds (83.4% were non-gapped contiguous sequences) with an N50 length of 57.6 kb (Supplementary Information). The assembly represented 97% of the 4.36-Gb genome as estimated by *K*-mer analysis (Supplementary Information). We also obtained 13,185 *Ae. tauschii* expressed sequence tag (EST) sequences using Sanger sequencing, of which 11,998 (91%) could be mapped to the scaffolds with more than 90% coverage (Supplementary Information).

To aid in gene identification, we performed RNA-Seq (53.2 Gb for a 117-Mb transcriptome assembly) on 23 libraries representing eight tissues including pistil, root, seed, spike, stamen, stem, leaf and sheath

(Supplementary Information). Using both evidence-based and *de novo* gene predictions, we identified 34,498 high-confidence protein-coding loci. FGENESH⁴ and GeneID models were supported by a 60% overlap with either our ESTs and RNA-Seq reads, or with homologous proteins. More than 76% of the gene models had a significant match (E value $\leq 10^{-5}$; alignment length $\geq 60\%$) in the GenBank non-redundant database. An additional 8,652 loci were predicted as low-confidence genes as a result of incomplete gene structure or limited expression data support (Supplementary Information). We also predicted a total of 2,505 transfer RNA, 358 ribosomal RNA, 35 small nuclear RNA and 78 small nucleolar RNA genes (Supplementary Information).

We found that more than 65.9% of the *Ae. tauschii* genome was composed of different transposable element (TE) families (Supplementary Information). About 5×10^6 Illumina reads of *Ae. tauschii* were mapped to hexaploid wheat repetitive sequences and we found that a comparable percentage of reads (more than 62.3%) could be classified as part of a TE sequence (Supplementary Fig. 6). This estimate is similar to that derived from a previous survey of Roche/454 sequences⁵. There were 410 different TE families, of which the 20 most abundant contributed more than 50% of the *Ae. tauschii* genome (Supplementary Table 9). A single peak of increased insertion activity was estimated to occur about 3–4 Myr ago by measuring the similarity of the assembled LTR retrotransposons (Supplementary Information), suggesting that the expansion of the *Ae. tauschii* genome was relatively recent and coincided with the abrupt climate change during the Pliocene Epoch⁶.

We constructed a high-density genetic map using an F₂ population of 490 individuals derived from a cross between the *Ae. tauschii* accessions Y2280 and AL8/78. The map, whose total length was 1059.8 centimorgans (cM), consisted of 151,083 single nucleotide polymorphism (SNP) markers developed by restriction-site-associated DNA (RAD) tag sequencing technology (Supplementary Fig. 13). Together with bin-mapped wheat ESTs⁷, SNPs and tags⁸, the genetic map was used to align 30,303 scaffolds (1.72 Gb; 30,697 genes) to chromosomes (Supplementary Information). The *Ae. tauschii* genes and scaffolds were also anchored to barley⁹ and *Brachypodium* chromosome maps¹⁰ (Fig. 1 and Supplementary Fig. 17). Calculation of K_a/K_s ratios (the ratio of non-synonymous substitutions to synonymous substitutions) for pairs of conserved orthologous genes showed that the average values between *Ae. tauschii* and barley (20,892 genes), *Brachypodium* (17,231 genes), rice (16,370 genes) and sorghum (18,623 genes) were 0.2214, 0.1888, 0.1736 and 0.1726, respectively,

¹National Key Facility for Crop Gene Resources and Genetic Improvement, Institute of Crop Science, Chinese Academy of Agricultural Sciences, Beijing 100081, China. ²BGI-Shenzhen, Shenzhen 518083, China. ³State Key Laboratory of Agrobiotechnology and School of Life Sciences, The Chinese University of Hong Kong, Shatin, New Territories, Hong Kong. ⁴Centre for Comparative Genomics, Murdoch University, Perth, WA 6150, Australia. ⁵MIPS/Institute for Bioinformatics and Systems Biology, Helmholtz Center Munich, D-85764 Neuherberg, Germany. ⁶Department of Vegetable Crops, College of Horticulture and Forestry, Huazhong Agricultural University, Wuhan 430070, China. ⁷Institute of Plant Biology, University of Zurich, CH-8008 Zurich, Switzerland. ⁸National Supercomputer Center in Tianjin, Tianjin 300457, China. ⁹International Maize and Wheat Improvement Center (CIMMYT), Texcoco CP 56130, Mexico. ¹⁰Department of Biology, University of Copenhagen, Copenhagen DK-2200, Denmark. ¹¹King Abdulaziz University, Jeddah 21589, Saudi Arabia.

*These authors contributed equally to this work.

†A list of participants and their affiliations appears in the Supplementary Information.

Table 1 | Overall statistics of sequencing and genome assembly

Assembly process	Library insert size (bp)	Read length (bp)	Effective data (Gb)	N50 (bp)	N50 number	Total length (Mb)	Gaps (Mb)
Contig assembly	167–764	44, 75, 100, 150	270	4,521	179,145	3,528	–
Scaffolding	2,000–20,000	44, 49, 90	128	58,011	19,405	4,244	1,122
Gap closure	167–764	44, 75, 100, 150	270	57,585	19,455	4,229	701
		114–263	65				
		~600*	18				

* Reads from 454 sequencing platform.

which indicated that most gene lineages evolved under purifying selection in *Ae. tauschii*. A total of 628 genes exhibited K_a/K_s ratios of more than 0.8 when compared with the other four species, indicating potential positive selection (innermost circle of Fig. 1). These genes were assigned to a wide range of molecular functions by using Gene Ontology (GO) analyses (Supplementary Table 14).

Ae. tauschii proteins were clustered with those of *Brachypodium*, rice, sorghum and barley (full-length complementary DNAs), and formed 23,202 orthologous groups (at least two members; Supplementary Information). In total, we identified 11,289 (barley/*Ae. tauschii*) and 14,675 (*Brachypodium*/*Ae. tauschii*) orthologous gene pairs. We found that 8,443 gene groups contained sequences from all five grass genomes, and 234 were specific to Pooideae (*Ae. tauschii*, *Brachypodium* and barley) and 587 were specific to Triticeae (*Ae. tauschii* and barley) (Fig. 2a). Enrichment analyses of both Pfam domains and GO terms showed that genes encoding NBS-LRR (nucleotide-binding-site

leucine-rich repeat) proteins were over-represented in *Ae. tauschii* relative to *Brachypodium* and rice^{11,12} (Supplementary Information). These observations are consistent with those reported in a recent study¹³. A total of 1,219 *Ae. tauschii* genes were similar to NBS-LRR genes (R gene analogues (RGAs))^{11,14} (Supplementary Information). This number is double that in rice (623) and sixfold that in maize (216)¹², indicating that the RGA family has substantially expanded in *Ae. tauschii*. We mapped 878 RGAs (72%) to specific positions across wheat chromosomes by using molecular marker–genome sequence alignment, which provides a large number of potential disease resistance loci for further investigation.

We found more genes for the cytochrome P450 family in *Ae. tauschii* (485) than in sorghum (365), rice (333), *Brachypodium* (262) or maize (261). This family of genes is important for abiotic stress response, especially in biosynthetic and detoxification pathways¹⁵. Using 178 manually curated cold-acclimation-related genes such

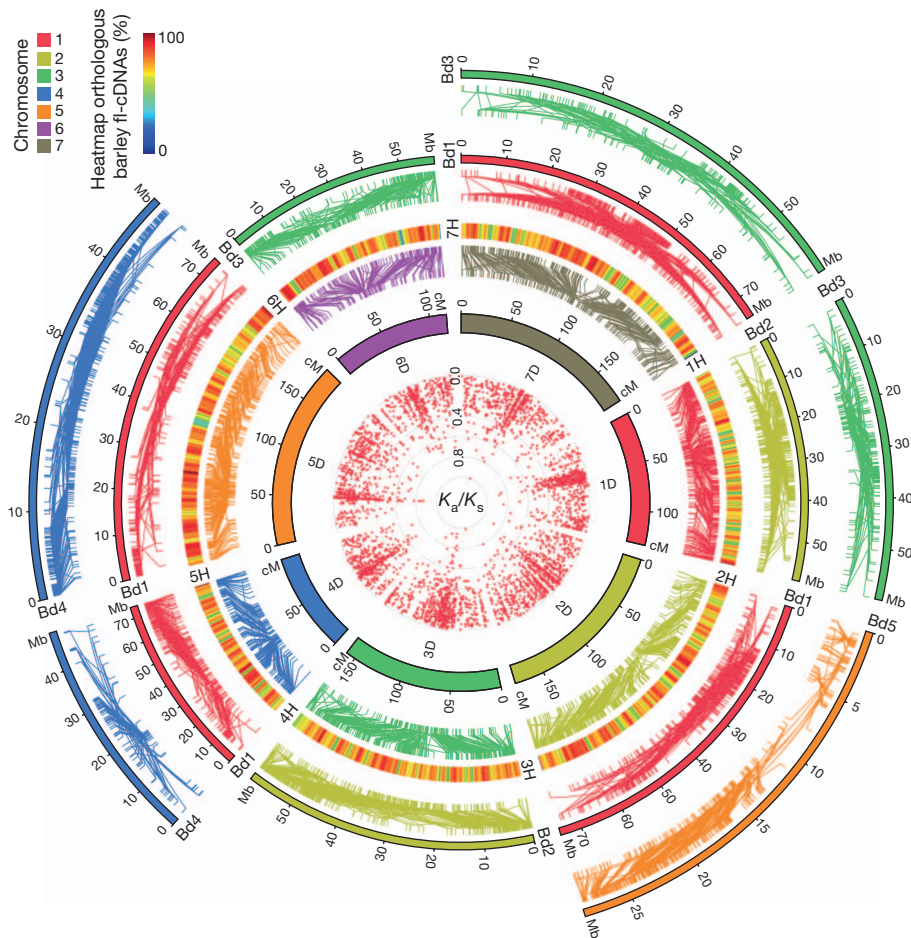


Figure 1 | Comparative analysis of *Ae. tauschii* ordered scaffolds versus barley and *Brachypodium*. The inner circle represents the seven *Ae. tauschii* chromosomes scaled according to the genetic map incorporating genome scaffolds. Red points show the K_a/K_s ratios between anchored *Ae. tauschii* genes and their putative orthologues in *Brachypodium*. Moving outwards, the second circle compares *Ae. tauschii* against the seven barley chromosomes⁹. The

heatmaps show the density distribution of barley cDNA loci that are aligned with *Ae. tauschii* genes. The outer two circles illustrate *Brachypodium* chromosomes according to conserved synteny with *Ae. tauschii*. The coloured lines below each chromosome identify putative orthologous gene pairs between *Ae. tauschii* genes, barley genes and *Brachypodium* genes.

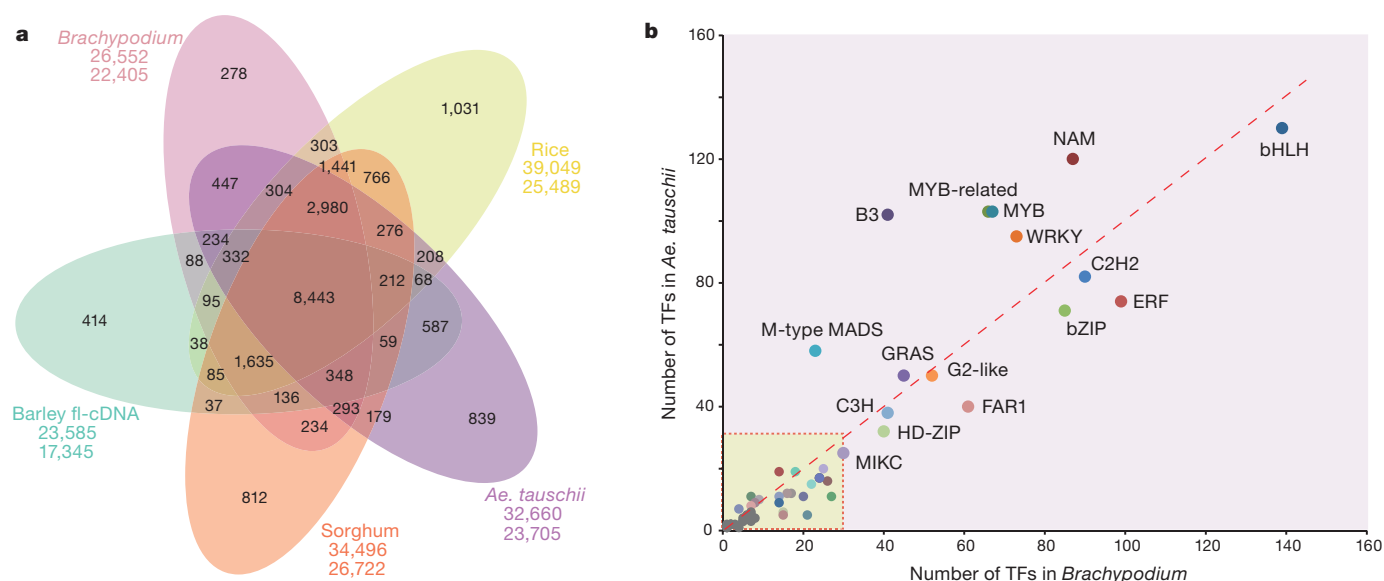


Figure 2 | *Ae. tauschii* gene families and transcription factors.

a, Distribution of orthologous gene families in *Ae. tauschii*, *Brachypodium*, sorghum, rice and barley. The number of gene families is represented in each intersection of the Venn diagram. The first number under the species name indicates the total number of genes annotated for a particular species, and the

second indicates the number of genes in groups for that organism. The difference between the two accounts for singleton genes that were not present in any cluster. **b**, The composition of transcription factors (TFs) in *Ae. tauschii* and *Brachypodium* composed of more than 30 members.

as the CCAAT-binding factor (CBF) transcription factors¹⁶, late-embryogenesis-abundant proteins (LEA) and osmoprotectant biosynthesis proteins (Supplementary Information) as queries, we identified 216 cold-related genes in the *Ae. tauschii* genome, in contrast to 164 genes in *Brachypodium*, 132 in rice, 159 in sorghum and 148 in maize. Some of these genes were specific to *Ae. tauschii* or Pooideae, including those encoding ice-recrystallization inhibition protein 1 precursor, DREB2 transcription factor α isoform and cold-responsive LEA/RAB-related COR protein. Expression analysis of RNA-Seq data showed that most of these *Ae. tauschii*-specific and Pooideae-specific genes were constitutively expressed in *Ae. tauschii* (Supplementary Fig. 23). In addition, 1,489 transcription factors (TFs) in 56 families were identified by using Pfam DNA-binding domains (Supplementary Information). *Ae. tauschii* had an excess of such TFs as MYB-related genes (103, in contrast with 66 in *Brachypodium* and 95 in maize), and these are also thought to be involved in various stress responses¹⁷. The M-type MADS-box genes (58, in contrast with 23 in *Brachypodium* and 34 in maize) are involved in regulation in plant reproduction¹⁸ (Fig. 2b and Supplementary Table 18). ARACNe¹⁹ co-expression analysis using RNA-Seq data predicted an expression network of 1,283 interactions (Supplementary Fig. 25), in which 13 TFs were associated with the expression of drought tolerance genes²⁰ (Supplementary Table 20).

We predicted a total of 159 (133 families) previously undescribed microRNAs (Supplementary Information), and identified segmental and tandem duplications in 42 members of the miR2118 family that were organized into two groups on 15 scaffolds (Supplementary Fig. 26). The miR399 family, which is involved in the regulation of inorganic phosphate homeostasis in rice²¹, was expanded (20 members in *Ae. tauschii*, compared with 11 in rice and 10 in maize), and may contribute to the ability of *Ae. tauschii* to grow in low-nutrient soils. The expansion of the miR2275 family (eight members in *Ae. tauschii*, compared with two in rice and four in maize) may contribute to the enhanced disease resistance of *Ae. tauschii* because phased short interfering RNAs initiated by miR2275 have been implicated in these activities²².

The *Ae. tauschii* genome served as the source for many grain quality genes in hexaploid wheat, creating a step improvement in the formation of the elastic dough essential for bread making². Grain quality genes include high-molecular-weight glutenin subunits (HMW-GS),

low-molecular-weight glutenin subunits (LMW-GS)²³, grain texture proteins (GSP; puroindolines)²⁴ and storage protein activator (SPA)²⁵. We identified two HMW-GS genes, five LMW-GS genes, one *Pina* gene, two *Pinb* genes, one GSP gene and one SPA gene in the *Ae. tauschii* genome sequence (Supplementary Information). As has been shown for the *Hardness* (*Ha*) locus²⁴, the GSP, *Pina* and *Pinb* genes were also organized in a cluster. RNA-Seq analysis showed that these grain quality genes were expressed predominantly in seeds (Supplementary Fig. 29).

The anchoring of more than 40% of the scaffold sequences to four genetic maps and to syntenic regions of other sequenced grass species provided a structural framework for integrating multiple maps by using shared markers (Fig. 1 and Supplementary Information). The co-localization of genes in scaffolds and genetically mapped quantitative trait loci (QTLs) will directly support map-based gene cloning. On chromosome 2D, for example, the locations of 33 QTLs or genes were integrated with scaffold information (<http://ccg.murdoch.edu.au/cmap/ccg-live/>) (Fig. 3 and Supplementary Information). Alignment of the *Ae. tauschii* genetic map with the wheat 2D consensus genetic map was unambiguous, with the exception of some single crossovers that were probably due to repetitive elements (dotted lines in Fig. 3). The genome sequence also provided the basis for the identification of more than 860,126 simple sequence repeats (SSRs), with trimers (37.7%) and tetramers (27.5%) as the most abundant SSR types (Supplementary Information). Together with the 711,907 SNPs identified by resequencing a roughly fivefold coverage of a second accession, Y2280 (Supplementary Information), the genomic resources reported here will promote map-based gene cloning and marker-assisted selection in wheat.

With its high base accuracy and nearly complete set of gene sequences, the *Ae. tauschii* draft genome sequence provides an essential reference for studying D genome diversity by re-sequencing additional accessions. Over the past half century, the introduction of new D genome diversity into synthetic wheat has been a major effort to expand bread wheat genetic diversity and to create environmentally resilient lines^{26,27}. The *Ae. tauschii* genome sequence should aid in identifying new elite alleles for agriculturally important traits to alleviate the worsening plight of global climate and environment changes²⁷.

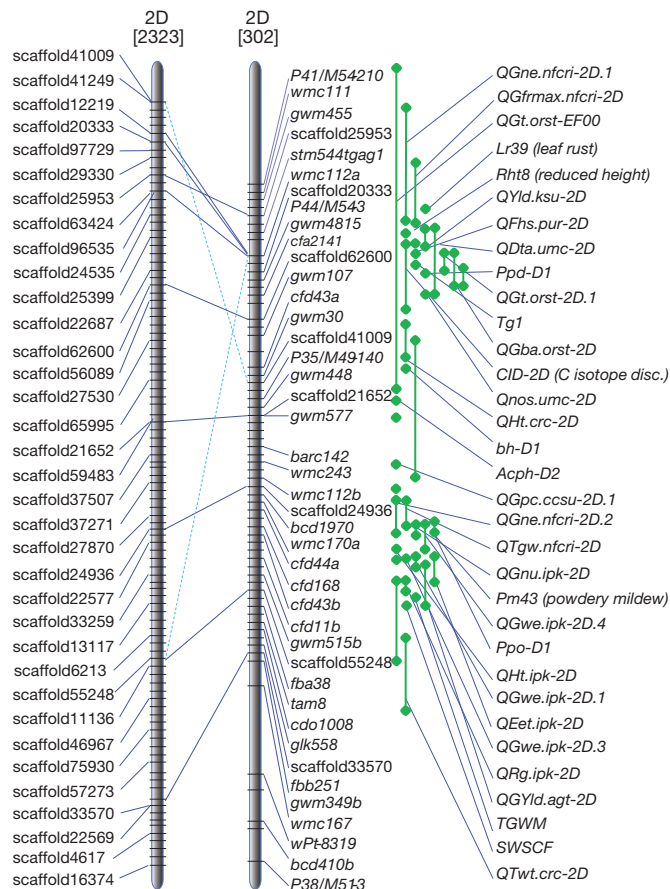


Figure 3 | An integrated genetic map of *Ae. tauschii* chromosome 2D. The *Ae. tauschii* genetic map was integrated with markers, scaffolds and mapped QTLs to assist in marker development and map-based cloning. Left: the *Ae. tauschii* molecular map used for synteny alignment in Fig. 1 was aligned to chromosome 2D (November 2011 consensus map, CMap; <http://ccg.murdoch.edu.au/cmap/ccg-live>) where sequence information was available. The original marker at a location is retained in CMap as a synonym. Right: within CMap, details for QTL locations are provided at a greater magnification to show all the markers in the regions of interest. The dotted lines indicate an ambiguous relationship that is most probably due to repetitive sequences.

METHODS SUMMARY

We selected *Ae. tauschii* ($2n = 14$) accession AL8/78 for sequencing. Plants were grown at 25 °C in a darkened chamber for two weeks; DNA was extracted from leaf tissue and purified with a standard phenol/chloroform extraction protocol. Sequencing libraries were constructed and sequenced on Illumina next-generation sequencing platforms (GAII and HiSeq2000). High-quality reads were assembled with SOAPdenovo³. Repeat sequences were identified by combining *de novo* approaches and sequence similarity at the nucleotide and protein levels. Gene models were predicted by combining homology-based, *de novo* and RNA-Seq-based methods. RNA-Seq reads were assembled with CAP3 (ref. 28) and CD-Hit²⁹ and were mapped to the draft genome with Tophat³⁰. See Supplementary Information for details and additional analyses.

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Supplementary Information is available in the online version of the paper.

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Author Contributions J.Z.J., Ji.W., Xu L., H.M.Y., Z.H.H., Lo.M., Ju.W., X.Y.K., X.Y.Z., R.L.J. and Y.Z.M. initiated the project and designed the study. X.Y.K., G.Y.Z., R.A., L.F.G., Qu.H., H.H.K., B.K., X.C.X., R.Z.Z. G.M.L. and C.M. performed the research. S.C.Z., Y.R.L., W.M.H., M.P., C.Z., D.L., C.G., M.S., K.M., Y.T., F.Y.Z., S.K.P., J.W.L., Q.S.L., Ji.C., C.Y.G., G.Y.H., Y.D.L., P.L., J.Y.W., J.Y.X. and J.L.G. generated and analysed the data. Q.J.X., H.F.Z. and Z.W.Q. developed the high-density genetic map. J.Z.J., S.C.Z., Lo.M., R.A., G.Y.Z., X.Y.K. and T.W. wrote the paper.

Author Information The genome sequence and the annotation are available from the National Centre for Biotechnology Information (NCBI) as BioProject ID PRJNA182898. This Whole Genome Shotgun project is deposited at DDBJ/EMBL/GenBank under accession number AOC000000000. The version described in this paper is the first version, AOC001000000. The Illumina sequencing reads are available in the Sequence Read Archive under accession number SRA030526, RNA-Seq sequences under SRA062662, and resequencing short reads under SRA063175. Genomic data are also available at the Comprehensive Library for Modern Biotechnology (CLiMB) repository under doi:10.5524/100054. Reprints and permissions information is

available at www.nature.com/reprints. The authors declare no competing financial interests. Readers are welcome to comment on the online version of the paper. Correspondence and requests for materials should be addressed to Ju.W. (wangj@genomics.cn), J.Z.J. (jzjia@mail.caas.net.cn), Lo.M. (maolong@caas.net.cn), Z.H.H. (zhhe@public3.bta.net.cn) and Xu L. (liuxu01@caas.cn).



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